What is claim is:

1. Particles for drug delivery by inhalation, comprising at least one active ingredient which is non-crystalline.

- 2. Particles according to Claim 1, further comprising a second or more active(s) and/or one or more excipients.
- 3. Particles for drug delivery according to Claim 1 or Claim 2, the particles containing a plurality of non-crystalline active ingredients.
- 4. Particles according to Claim 3, further **comprising** an outer surface of the particles being substantially smooth.
- 5. Particles according to Claim 3, particles being substantially spherical.
- 6. Particles according to Claim 5, particles being oblate spheroidal.
- 7. Particles according to Claim 3, particles being substantially oval.
- 8. Particles according to Claim 3, particles being substantially elliptical.
- 9. Particles according to any preceding claim, having a particle size in the range $0.5 \mu m$ to $5 \mu m$.
- 10. Particles according to Claim 9, the particle size being between 1 μm to 3 μm.
- 11. Particles according to Claim 9 when dependent on Claim 7 or Claim 8, the longer axis of an oval or elliptical particle having a length between 1 µm to 3 µm.
- 12. Particles according to any preceding Claim, the particles being electrically uncharged.
- 13. Particles according to any preceding claim, provided by a method selected from the group comprising rapid expansion of supercritical solutions, precipitation from gas saturated solutions, gas anti-solvent systems, aerosol solvent extraction systems and spray drying processes.
- 14. Particles according to any preceding claim, there being from two-to four active ingredients.

 Particles according to any preceding claim, comprising a pharmaceutically acceptable particular excipient or excipients.

- 16. Particles according to any preceding claim, the active ingredients comprising a £2 agonist and a steroid.
- 17. Particles according to Claim 15, comprising fluticasone-dipropionate and salmeterol xinafoate.
- 18. Particles according to any one of Claims 1 to 15, comprising formoterol and budesonide.
- 19. Particles for drug delivery according to any of claims 2 to 18, the or each excipient being soluble in conditions obtaining in the nose, lung(s) or mouth of a human or animal.
- 20. An inhalation composition, comprising particles which incorporate at least one active ingredient which is non-crystalline.
- 21. A composition according to Claim 20, further comprising a second or more active(s) and/or one or more excipients.
- 22. A composition according to Claim 20 or 21, the particles containing a plurality of active ingredients, which active ingredients are non-crystalline.
- 23. A composition according to Claims 20 to 22, there being from two to four active ingredients.
- 24. A composition according to any of Claims 21 to 23, the particles comprising a pharmaceutically acceptable excipient within the particle.
- 25. A composition according to any of Claims 21 to 24, the particles comprising a pharmaceutically acceptable excipient or excipients where a main excipient is in a greater proportion than the active or actives.
- 26. A composition according to Claim 25, the main excipient being Mannitol or PVP.
- 27. A composition according to any of Claims 21 to 26, comprising one or more additional carrier excipient(s).

28. A composition according to Claim 27, said excipient(s) comprising a modifier or stabiliser.

- 29. A composition according to Claim 27, said excipient(s) comprising a chemical buffer, antioxidant and the like.
- 30. A composition according to Claim 27, said excipient(s) comprising a surface modifier or surfactant.
- 31. A composition according to any of Claims 20 to 30, the outer surface of the particles being substantially smooth.
- 32. A composition according to Claim 31, the particles being substantially spherical.
- 33. A composition according to Claim 32, the particles being oblate spheroidal.
- 34. A composition according to Claim 31, the particles being substantially oval.
- 35. A composition according to Claim 31, the particles being substantially elliptical.
- 36. A composition according to any of Claims 20 to 35, the particles thereof having a particle size in the range 0.5μm 5μm.
- 37. A composition according to Claim 36, having a particle size of 1μ to 3μ.
- 38. A composition according to Claim 37, the particles being electrically uncharged.
- 39. A composition according to any of Claims 20 to 38, provided by a method selected from the group comprising rapid expansion of supercritical solutions, precipitation from gas saturated solutions, gas anti-solvent systems, aerosol solvent extraction systems, and a spray drying process.
- 40. A composition according to any of claims 20 to 39, comprising fluticasone and salmeterol xinafoate as active ingredients.
- 41. A composition according to any of Claims 20 to 39, the particles comprising formoterol and budesonide as active ingredients.
- 42. A composition according to any of Claims 29 to 39, the particles containing one or more cannabinoids as an active ingredient.
- 43. A composition according to claim 42, the cannabinoid comprising delta-8 or delta-9 tetrahydrocannabinol.

44. An inhaler device, comprising an inhalation composition according to any of Claims 20 to 43.

- 45. A pulmonary nasal inhalation device, comprising an inhalation composition according to any of Claims 20 to 44.
- 46. A device according to Claim 44 or Claim 45, the main excipient being non-soluble in the propellant or propellants.